

FILE 'MEDLINE' ENTERED AT 16:10:24 ON 08 SEP 2004

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=> e REID IAN R/au

E1	4	REID IAN MAITLAND/AU
E2	6	REID IAN N/AU
E3	156 -->	REID IAN R/AU
E4	13	REID IAN REGINALD/AU
E5	6	REID III G L/AU
E6	2	REID III G R/AU
E7	1	REID III T/AU
E8	8	REID III T J/AU
E9	3	REID IONA/AU
E10	2	REID IONA M/AU
E11	1	REID ISLAY S/AU
E12	2	REID IV D S/AU

=> s l3 and calcium

L3 NOT FOUND

The L-number entered could not be found. To see the definition
of L-numbers, enter DISPLAY HISTORY at an arrow prompt (=>).

=> s e3 and calcium

L1 47 "REID IAN R"/AU AND CALCIUM

=> s l1 and (cholesterol or lipid? or hdl or ldl)

L2 7 L1 AND (CHOLESTEROL OR LIPID? OR HDL OR LDL)

=> dup rem l2

PROCESSING COMPLETED FOR L2

L3 3 DUP REM L2 (4 DUPLICATES REMOVED)

=> d ibib abs 1-3

L3	ANSWER 1 OF 3	MEDLINE on STN	DUPLICATE 1
ACCESSION NUMBER:	2004018286	MEDLINE	
DOCUMENT NUMBER:	PubMed ID: 14715041		
TITLE:	Effects of calcium supplementation on circulating lipids : potential pharmacoeconomic implications.		
AUTHOR:	Reid Ian R		
CORPORATE SOURCE:	Department of Medicine, University of Auckland, Auckland, New Zealand.. i.reid@auckland.ac.nz		
SOURCE:	Drugs & aging, (2004) 21 (1) 7-17. Ref: 34 Journal code: 9102074. ISSN: 1170-229X.		
PUB. COUNTRY:	New Zealand		
DOCUMENT TYPE:	Journal; Article; (JOURNAL ARTICLE) General Review; (REVIEW) (REVIEW, TUTORIAL)		
LANGUAGE:	English		
FILE SEGMENT:	Priority Journals		
ENTRY MONTH:	200404		
ENTRY DATE:	Entered STN: 20040113 Last Updated on STN: 20040413 Entered Medline: 20040412		

AB For about a century there has been recognition that **calcium** and **lipids** bind to one another in the gut, each interfering with the other's absorption. **Calcium** also causes malabsorption of bile acids, which is likely to contribute further to malabsorption of fat. High dietary **calcium** intakes may also have stimulatory effects on lipolysis. These mechanisms provide a basis for hypothesising that **calcium** supplementation may impact on circulating **lipid** concentrations, and there is now a significant amount of observational and trial data indicating that this is the case. The largest randomised controlled trial of **calcium** effects on **lipids** was carried out in 223 healthy postmenopausal women, and found that low density lipoprotein-**cholesterol** (LDL-C) decreased 6.3% and high density lipoprotein-**cholesterol** (HDL-C) increased by 7.3% at 1-year. The resultant 16.4% increase in HDL-C/LDL-C ratio would be predicted to reduce cardiovascular event rates by 20-30%, which is consistent with the available observational data. There are no trial data addressing this question and it is possible that other **lipid**-lowering agents, such as hydroxymethylglutaryl coenzyme A reductase inhibitors, might impact on cardiac event rates by mechanisms other than by lowering **cholesterol** levels. Therefore, caution is appropriate in incorporating these findings into clinical practice, but the balance of evidence suggests that **calcium** is a cost-effective adjunct to the dietary management of hyperlipidaemia.

L3 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 2002:696671 CAPLUS
 DOCUMENT NUMBER: 137:216323
 TITLE: Method of administering **calcium** citrate
 INVENTOR(S): Reid, Ian R.
 PATENT ASSIGNEE(S): Uniservices Ltd., N. Z.
 SOURCE: U.S. Pat. Appl. Publ., 13 pp.
 CODEN: USXXCO
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2002128320	A1	20020912	US 2001-16371	20011210
WO 2003049668	A2	20030619	WO 2002-IB5759	20021210
WO 2003049668	A3	20040617		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

PRIORITY APPLN. INFO.: US 2001-16371 A 20011210

AB A method of increasing a high-d. lipoprotein level in plasma of a postmenopausal woman by administering a pharmaceutical formulation containing **calcium** citrate is described. The therapeutically ED of **calcium** citrate is equivalent to at least about 1 g elemental **calcium**. An oral pharmaceutical composition and a dietary supplement comprises **calcium** citrate in an amount sufficient to provide about 10 mg to about 1 g elemental **calcium** to a diet of a postmenopausal woman.

L3 ANSWER 3 OF 3 MEDLINE on STN DUPLICATE 2
 ACCESSION NUMBER: 2002171646 MEDLINE
 DOCUMENT NUMBER: PubMed ID: 11904107
 TITLE: Effects of **calcium** supplementation on serum

lipid concentrations in normal older women: a randomized controlled trial.
COMMENT: Comment in: Am J Med. 2003 May;114(7):620-1; author reply 621. PubMed ID: 12753892
AUTHOR: **Reid Ian R**; Mason Barbara; Horne Anne; Ames Ruth; Clearwater Judith; Bava Usha; Orr-Walker Brandon; Wu Fiona; Evans Margaret C; Gamble Gregory D
CORPORATE SOURCE: Department of Medicine, University of Auckland, Auckland, New Zealand.
SOURCE: American journal of medicine, (2002 Apr 1) 112 (5) 343-7. Journal code: 0267200. ISSN: 0002-9343.
PUB. COUNTRY: United States
DOCUMENT TYPE: (CLINICAL TRIAL)
Journal; Article; (JOURNAL ARTICLE)
(RANDOMIZED CONTROLLED TRIAL)
LANGUAGE: English
FILE SEGMENT: Abridged Index Medicus Journals; Priority Journals
ENTRY MONTH: 200204
ENTRY DATE: Entered STN: 20020321
Last Updated on STN: 20020404
Entered Medline: 20020402

AB PURPOSE: To determine the effect of supplementation with **calcium** citrate on circulating **lipid** concentrations in normal older women. SUBJECTS AND METHODS: As part of a study of the effects of **calcium** supplementation on fractures, we randomly assigned 223 postmenopausal women (mean [\pm SD] age, 72 \pm 4 years), who were not receiving therapy for hyperlipidemia or osteoporosis, to receive **calcium** (1 g/d, n = 111) or placebo (n = 112) for 1 year. Fasting serum **lipid** concentrations, including high-density lipoprotein (HDL) **cholesterol** and low-density lipoprotein (LDL) **cholesterol**, were obtained at baseline, and at 2, 6, and 12 months. RESULTS: After 12 months, HDL **cholesterol** levels and the HDL **cholesterol** to LDL **cholesterol** ratio had increased more in the **calcium** group than in the placebo group (mean between-group differences in change from baseline: for HDL **cholesterol**, 0.09 mmol/L (95% confidence interval [CI]: 0.02 to 0.17; P = 0.01); for HDL/LDL **cholesterol** ratio, 0.05 (95% CI: 0.02 to 0.08; P = 0.001). This was largely due to a 7% increase in HDL **cholesterol** levels in the **calcium** group, with a nonsignificant 6% decline in LDL **cholesterol** levels. There was no significant treatment effect on triglyceride level (P = 0.48). CONCLUSION: **Calcium** citrate supplementation causes beneficial changes in circulating **lipids** in postmenopausal women. This suggests that a reappraisal of the indications for **calcium** supplementation is necessary, and that its cost effectiveness may have been underestimated.